

## AMENDMENTS TO THE CLAIMS

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Claim 1 (currently amended). A method of screening ~~proteins or and~~ polypeptides to identify a protein or polypeptide having a biological activity of interest, which comprises the sequential steps of (i) forming a gene first library of polynucleotide clones; and synthesizing individual proteins, which can then be screened (ii) expressing an individual protein or polypeptide from each clone in the first library to form a second library of individual proteins and polypeptides therefrom; (iii) assaying the second library to select an individual protein or polypeptide in the second library having a biological activity of interest; and (iv) identifying the protein or polypeptide selected in step (iii) by sequencing the polynucleotide from the first library that encodes the selected protein or polypeptide.

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Claim 2 (currently amended). A method as claimed in claim 1 wherein the individual proteins ~~or and~~ polypeptides ~~can be screened for (a) in the second library are assayed for a biological activity selected from the group consisting of an enzymatic protein or polypeptide modification, and/or (b) binding to one or more other another molecule, molecules/ligands and/or (c) binding or biological activity on cells or tissues to a cell or tissue, and modulating the metabolism of a cell or tissue.~~

Claim 3 (currently amended). A method as claimed in claim 2 wherein the ~~gene~~ first library is derived from a library of cellular mRNA from one or more cells or tissues.

Claim 4 (currently amended). A method as claimed in claim 2 wherein the ~~gene library encodes proteins or polypeptides comprising a library of variable molecules, such as~~ second library of individual proteins and polypeptides comprises fragments of antibody variable regions.

Claim 5 (currently amended). A method as claimed in claim 4 wherein the ~~proteins or polypeptides are screened for~~ biological activity of interest is binding to one or more proteins or polypeptides from a cell or tissue.

Claim 6 (withdrawn).

Claim 7 (currently amended). A method as claimed in claim 1 wherein the ~~individual members of the gene library are initially~~ first library of polynucleotide clones is

distributed into ~~one or more arrays whereby each gene is then~~ an array of polynucleotides, and in step (ii) each polynucleotide in the array is then expressed to generate ~~one or more protein or polypeptide arrays~~ an array of individual proteins and polypeptides.

Claim 8 (currently amended). A method as claimed in claim 7 wherein the array of individual proteins ~~or~~ and polypeptides ~~are~~ is immobilized onto a solid phase.

Claim 9 (withdrawn).

Claim 10 (currently amended). A method as claimed in claim 8 wherein the solid phase is a ~~continuous surface such as a glass plate~~ and wherein the proteins ~~or~~ and polypeptides are immobilized at specific loci on the surface of the plate.

Claim 11 (currently amended). A method as claimed in claim 7 wherein the individual proteins ~~or~~ and polypeptides are expressed from the polynucleotides in the first library by *in vitro* transcription and translation.

Claims 12-13 (cancelled).

Claim 14 (withdrawn).

Claim 15 (currently amended). A method for screening proteins ~~or~~ and polypeptides to identify a protein or polypeptide having a biological activity of interest, which comprises the sequential steps of:

- (i) ~~Generating a gene~~ generating a first library of polynucleotides in the form of clones selected from the group consisting of DNA molecules, RNA molecules, cell colonies, or and plaques;
- (ii) ~~Converting the nucleic acid~~ expressing a polynucleotide from each clone in the first library using *in vitro* translation to generate ~~proteins or polypeptides~~ a second library of individual proteins and polypeptides therefrom;
- (iii) ~~Dispensing aliquots~~ dispensing an aliquot of each protein or polypeptide in the second library into a specific loci in a multi-well plate or a solid phase to form a protein or polypeptide array; and
- (iv) ~~Bringing the arrays generated (iii) into contact with one or more extracts from cells or tissues or with one or more cells or tissues per se~~

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~~in order to screen for protein or polypeptide modification or for binding to one or more molecules from the one or more extracts contacting the array generated in step (iii) with a material selected from the group consisting of a cell extract, a tissue extract, a cell sample, and a tissue sample;~~

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- ~~(v) assaying each protein and polypeptide in the array to select an individual protein or polypeptide that interacts with the material contacting the array in step (iv), and~~
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- ~~(vi) identifying the individual protein or polypeptide selected in step (v) by sequencing the polynucleotide that encodes the selected protein or polypeptide;~~

~~wherein the interaction of the protein or polypeptide with the material contacting the array in step (v) is an interaction selected from the group consisting of modification of a protein or polypeptide in the array, binding of a protein or polypeptide in the array to a molecule from a cell, and binding of a protein or polypeptide in the array to a molecule from a tissue.~~

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Claims 16-25 (cancelled).

Claims 26-42 (withdrawn).

Claims 43-56 (cancelled).

Claim 57 (withdrawn).